

Requested Patent: WO2005048948A2

Title: UREA DERIVATIVES AS KINASE MODULATORS ;

Abstracted Patent: WO2005048948 ;

Publication Date: 2005-06-02 ;

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Application Number: WO2004US38288 20041115 ;

Priority Number(s):

US20030520273P 20031113; US20030527094P 20031203; US20030531082P 20031218; US20030531243P 20031218 ;

IPC Classification: A61K ;

Equivalents: WO2005048953 ;

ABSTRACT:

The invention provides methods and compositions for treating conditions mediated by various kinases wherein derivatives of urea compounds are employed. The invention also provides methods of using the compounds and/or compositions in the treatment of a variety of diseases and unwanted conditions in subjects.

(19) World Intellectual Property
Organization
International Bureau



(43) International Publication Date
2 June 2005 (02.06.2005)

PCT

(10) International Publication Number
WO 2005/048948 A2

- (51) International Patent Classification⁷: **A61K** (US). LOCKHART, David, J. [US/US]; 510 Torrey Point Road, Del Mar, CA 92014 (US).
- (21) International Application Number: PCT/US2004/038288 (74) Agents: HADDACH, Aubrey, A. et al.; Wilson Sonsini Goodrich & Rosati, 650 Page Mill Road, Palo Alto, CA 94306-1050 (US).
- (22) International Filing Date: 15 November 2004 (15.11.2004) (81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.
- (25) Filing Language: English
- (26) Publication Language: English
- (30) Priority Data:
60/520,273 13 November 2003 (13.11.2003) US
60/527,094 3 December 2003 (03.12.2003) US
60/531,082 18 December 2003 (18.12.2003) US
60/531,243 18 December 2003 (18.12.2003) US
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- (84) Designated States (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).
- Published:
— without international search report and to be republished upon receipt of that report
- For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: UREA DERIVATIVES AS KINASE MODULATORS

(57) Abstract: The invention provides methods and compositions for treating conditions mediated by various kinases wherein derivatives of urea compounds are employed. The invention also provides methods of using the compounds and/or compositions in the treatment of a variety of diseases and unwanted conditions in subjects.

WO 2005/048948 A2

UREA DERIVATIVES AS KINASE MODULATORS

This application claims priority to US Provisional Application No. 60/520,273, filed November 13, 2003, US Provisional Application No. 60/527,094, filed December 3, 2003, US Provisional Application No. 60/531,243, filed December 18, 2003, and US Provisional Application No. 60/531,082, filed December 18, 2003, the contents of which are
5 incorporated herein by reference in their entirety.

BACKGROUND

Protein kinases (PKs) play a role in signal transduction pathways regulating a number of cellular functions, such as cell growth, differentiation, and cell death. PKs are enzymes that
10 catalyze the phosphorylation of hydroxy groups on tyrosine, serine and threonine residues of proteins, and can be conveniently broken down into two classes, the protein tyrosine kinases (PTKs) and the serine-threonine kinases (STKs). Growth factor receptors with PTK activity are known as receptor tyrosine kinases. Protein receptor tyrosine kinases are a family of tightly regulated enzymes, and the aberrant activation of various members of the family is
15 one of the hallmarks of cancer. The protein-tyrosine kinase family, which includes Bcr-Abl tyrosine kinase, can be divided into subgroups that have similar structural organization and sequence similarity within the kinase domain. The members of the type III group of receptor tyrosine kinases include the platelet-derived growth factor (PDGF) receptors (PDGF receptors α and β), colony-stimulating factor (CSF-1) receptor (CSF-1R, c-Fms), FLT-3, and
20 stem cell or steel factor receptor (c-kit). A more complete listing of the known Protein receptor tyrosine kinases subfamilies is described in Plowman et al., DN&P, 7(6):334-339 (1994), which is incorporated by reference, including any drawings, as if fully set forth herein. Furthermore, for a more detailed discussion of "non-receptor tyrosine kinases", see Bolen, Oncogene, 8:2025-2031 (1993), which is incorporated by reference, including any
25 drawings, as if fully set forth herein.

Hematologic cancers, also known as hematologic or hematopoietic malignancies, are cancers of the blood or bone marrow; including leukemia and lymphoma. Acute myelogenous leukemia (AML) is a clonal hematopoietic stem cell leukemia that represents ~90% of all acute leukemias in adults. See e.g., Lowenberg et al., *N. Eng. J. Med.* 341:1051-
30 62 (1999). While chemotherapy can result in complete remissions, the long term disease-free survival rate for AML is about 14% with about 7,400 deaths from AML each year in the

United States. The single most commonly mutated gene in AML is FLT3 kinase. See e.g., Abu-Duhier et al., *Br. J. Haematol.* 111:190-05 (2000); Kiyoi et al., *Blood* 93:3074-80 (1999); Kottaridis et al., *Blood* 98:1752-59 (2001); Stirewalt et al., *Blood* 97:3589-95 (2001). Such mutations also indicate a poor prognosis for the patient.

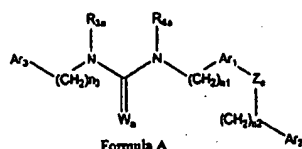
5 The compounds provided by the present invention are urea derivatives of substituted aryls and hetroaryls, e.g., isoxazoles, pyrazoles and isothiazoles. Urea derivatives of pyrazoles have been reported to be selective p38 kinase inhibitors by Dumas, J., et al., *Bioorg. Medic. Chem. Lett.* 10:2051-2054 (2000). Oxazoles and isopyrazoles are suggested as blockers of cytokine production in WO 00/43384 published 27 July 2000. Urea derivatives of isoxazole and pyrazoles are described as inhibitors of RAF kinase in WO 99/32106 published 1 July 1999. Such compounds are also described as p38 kinase inhibitors by Dumas, J., et al., *Bioorg. Medic. Chem. Lett.* 10:2047-2050 (2000). These compounds are also suggested as p38 kinase inhibitors in PCT publication WO 99/32111 published 1 July 1999.

15 There remains a need for additional compounds that are effective in inhibiting kinase activity. Given the complexities of signal transduction with the redundancy and crosstalk between various pathways, the identification of specific kinase inhibitors permits accurate targeting with limited inhibition of other pathways, thus reducing the toxicity of such inhibitory compounds.

20 SUMMARY OF THE INVENTION

The present invention provides compounds which modulate kinase activity, and in some embodiments inhibit protein tyrosine kinases or a specific kinase or kinase class. In some embodiments, the compositions and methods for treating and preventing conditions and diseases, such as cancer, hematologic malignancies, cardiovascular disease, inflammation or multiple sclerosis. The compounds of the invention can be delivered alone or in combination with additional agents, and are used for the treatment and/or prevention of conditions and diseases. As used throughout the specification, unless otherwise stated, each of the substituents is as previously defined.

30 Provided herein are compositions and methods for treating a disease comprising administering to a subject in need thereof an effective amount of a kinase modulating compound having the structure:



wherein:

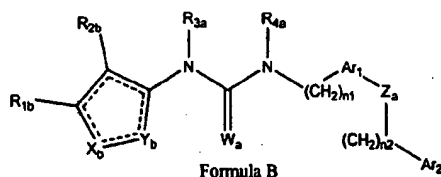
- (a) R_{3a} and R_{4a} are each a suitable substituent independently selected from hydrogen, or an alkyl, alkenyl, heteroalkyl, haloalkyl, alkynyl, aryl, cycloalkyl, heterocycloalkyl, heteroaryl group unsubstituted or substituted with one or more suitable substituents independently selected from the group consisting of: halogens; -CN; and -NO₂; and alkyl, alkenyl, heteroalkyl, haloalkyl, alkynyl, aryl, cycloalkyl, heterocycloalkyl, heteroaryl, -(CH₂)_zCN where z is a whole integer, preferably from 0 to 4, =NH, -NHOH, -OH, -C(O)H, -OC(O)H, -C(O)OH, -OC(O)OH, -OC(O)OC(O)H, -OOH, -C(NH)NH₂, -NHC(NH)NH₂, -C(S)NH₂, -NHC(S)NH₂, -NHC(O)NH₂, -S(O₂)H, -S(O)H, -NH₂, -C(O)NH₂, -OC(O)NH₂, -NHC(O)H, -NHC(O)OH, -C(O)NHC(O)H, -OS(O₂)H, -OS(O)H, -OSH, -SC(O)H, -S(O)C(O)OH, -SO₂C(O)OH, -NHSH, -NHS(O)H, -NH₂SO₂H, -C(O)SH, -C(O)S(O)H, -C(O)S(O₂)H, -C(S)H, -C(S)OH, -C(SO)OH, -C(SO₂)OH, -NHC(S)H, -OC(S)H, -OC(S)OH, -OC(SO₂)H, -S(O₂)NH₂, -S(O)NH₂, -SNH₂, -NHCS(O₂)H, -NHC(SO)H, -NHC(S)H, and -SH groups unsubstituted or substituted with one or more suitable substituents independently selected from the group consisting of halogens, =O, -NO₂, -CN, -(CH₂)_z-CN where z is a whole integer, preferably from 0 to 4, -OR_c, -NR_cOR_c, -NR_cR_c, -, C(O)NR_c, -C(O)OR_c, -C(O)R_c, -NR_cC(O)NR_c, -NR_cC(O)R_c, -OC(O)OR_c, -OC(O)NR_cR_c, -SR_c, unsubstituted alkyl, unsubstituted alkenyl, unsubstituted alkynyl, unsubstituted aryl, unsubstituted cycloalkyl, unsubstituted heterocycloalkyl, and unsubstituted heteroaryl, or two or more substituents cyclize to form a fused or spiro polycyclic cycloalkyl, heterocycloalkyl, aryl, or heteroaryl group, where each R_c is independently selected from hydrogen, unsubstituted alkyl, unsubstituted alkenyl, unsubstituted alkynyl, unsubstituted aryl, unsubstituted cycloalkyl, unsubstituted heterocycloalkyl, and unsubstituted heteroaryl, or two or more R_c groups together cyclize to form part of a heteroaryl or heterocycloalkyl group unsubstituted or substituted with an unsubstituted alkyl group; or where R_{3a} and R_{4a} together cyclize to form part of a heteroaryl or heterocycloalkyl group unsubstituted or substituted with one or more suitable substituents selected from

halogen, =O; =S; -CN; -NO₂, or an alkyl, alkenyl, heteroalkyl, haloalkyl, alkynyl, aryl, cycloalkyl, heterocycloalkyl, heteroaryl group unsubstituted or substituted with one or more suitable substituents independently selected from the group consisting of: halogens; =O; =S; -CN; and -NO₂; and alkyl, alkenyl, heteroalkyl, haloalkyl, alkynyl, aryl, cycloalkyl, heterocycloalkyl, heteroaryl, -(CH₂)_zCN where z is a whole integer, preferably from 0 to 4, =NH, -NHOH, -OH, -C(O)H, -OC(O)H, -C(O)OH, -OC(O)OH, -OC(O)OC(O)H, -OOH, -C(NH)NH₂, -NHC(NH)NH₂, -C(S)NH₂, -NHC(S)NH₂, -NHC(O)NH₂, -S(O₂)H, -S(O)H, -NH₂, -C(O)NH₂, -OC(O)NH₂, -NHC(O)H, -NHC(O)OH, -C(O)NHC(O)H, -OS(O₂)H, -OS(O)H, -OSH, -SC(O)H, -S(O)C(O)OH, -SO₂C(O)OH, -NHSH, -NHS(O)H, -NHSO₂H, -C(O)SH, -C(O)S(O)H, -C(O)S(O₂)H, -C(S)H, -C(S)OH, -C(SO)OH, -C(SO₂)OH, -NHC(S)H, -OC(S)H, -OC(S)OH, -OC(SO₂)H, -S(O₂)NH₂, -S(O)NH₂, -SNH₂, -NHCS(O₂)H, -NHC(SO)H, -NHC(S)H, and -SH groups unsubstituted or substituted with one or more suitable substituents independently selected from the group consisting of halogens, =O, -NO₂, -CN, -(CH₂)_z-CN where z is a whole integer, preferably from 0 to 4, -OR_c, -NR_cOR_c, -NR_cR_c, -C(O)NR_c, -C(O)OR_c, -C(O)R_c, -NR_cC(O)NR_cR_c, -NR_cC(O)R_c, -OC(O)OR_c, -OC(O)NR_cR_c, -SR_c, unsubstituted alkyl, unsubstituted alkenyl, unsubstituted alkynyl, unsubstituted aryl, unsubstituted cycloalkyl, unsubstituted heterocycloalkyl, and unsubstituted heteroaryl, or two or more substituents cyclize to form a fused or spiro polycyclic cycloalkyl, heterocycloalkyl, aryl, or heteroaryl group, where each R_c is independently selected from hydrogen, unsubstituted alkyl, unsubstituted alkenyl, unsubstituted alkynyl, unsubstituted aryl, unsubstituted cycloalkyl, unsubstituted heterocycloalkyl, and unsubstituted heteroaryl, or two or more R_c groups together cyclize to form part of a heteroaryl or heterocycloalkyl group unsubstituted or substituted with an unsubstituted alkyl group;

(b) Ar₁, Ar₂ and Ar₃ are each independently an aryl, heteroaryl, cycloalkyl or heterocycloalkyl group unsubstituted or substituted with one or more suitable substituents independently selected from the group consisting of: halogens; =O; =S; -CN; and -NO₂; and alkyl, alkenyl, heteroalkyl, haloalkyl, alkynyl, aryl, cycloalkyl, heterocycloalkyl, heteroaryl, -(CH₂)_zCN where z is a whole integer, preferably from 0 to 4, =NH, -NHOH, -OH, -C(O)H, -OC(O)H, -C(O)OH, -OC(O)OH, -OC(O)OC(O)H, -OOH, -C(NH)NH₂, -NHC(NH)NH₂, -C(S)NH₂, -NHC(S)NH₂, -NHC(O)NH₂, -S(O₂)H,

- S(O)H, -NH₂, -C(O)NH₂, -OC(O)NH₂, -NHC(O)H, -NHC(O)OH, -C(O)NHC(O)H,
 -OS(O₂)H, -OS(O)H, -OSH, -SC(O)H, -S(O)C(O)OH, -SO₂C(O)OH, -NHSH,
 -NHS(O)H, -NHSO₂H, -C(O)SH, -C(O)S(O)H, -C(O)S(O₂)H, -C(S)H, -C(S)OH,
 -C(SO)OH, -C(SO₂)OH, -NHC(S)H, -OC(S)H, -OC(S)OH, -OC(SO₂)H, -S(O₂)NH₂,
 5 -S(O)NH₂, -SNH₂, -NHCS(O₂)H, -NHC(SO)H, -NHC(S)H, and -SH groups
 unsubstituted or substituted with one or more suitable substituents independently selected
 from the group consisting of halogens, =O, -NO₂, -CN, -(CH₂)_z-CN where z is a whole
 integer, preferably from 0 to 4, -OR_c, -NR_cOR_c, -NR_cR_c, -C(O)NR_c, -C(O)OR_c,
 -C(O)R_c, -NR_cC(O)NR_cR_c, -NR_cC(O)R_c, -OC(O)OR_c, -OC(O)NR_cR_c, -SR_c, unsubstituted
 10 alkyl, unsubstituted alkenyl, unsubstituted alkynyl, unsubstituted aryl, unsubstituted
 cycloalkyl, unsubstituted heterocycloalkyl, and unsubstituted heteroaryl, or two or more
 substituents cyclize to form a fused or spiro polycyclic cycloalkyl, heterocycloalkyl, aryl,
 or heteroaryl group, where each R_c is independently selected from hydrogen,
 unsubstituted alkyl, unsubstituted alkenyl, unsubstituted alkynyl, unsubstituted aryl,
 15 unsubstituted cycloalkyl, unsubstituted heterocycloalkyl, and unsubstituted heteroaryl, or
 two or more R_c groups together cyclize to form part of a heteroaryl or heterocycloalkyl
 group unsubstituted or substituted with an unsubstituted alkyl group;
 (c) n₁ is 0, 1, 2, 3 or 4;
 (d) n₂ is 0, 1, 2, 3 or 4;
 20 (e) n₃ is 0, 1, 2, 3 or 4;
 (f) Z_a is a bond or is selected from S, O, N, NR_c, C(O)NR_c, NR_cC(O), and CR_c,
 wherein R_c is a suitable substituent selected from hydrogen, unsubstituted alkyl,
 unsubstituted alkenyl, unsubstituted alkynyl, unsubstituted aryl, unsubstituted cycloalkyl,
 unsubstituted heterocycloalkyl, or unsubstituted heteroaryl group; and
 25 (g) W_a is S or O;
 or a pharmaceutically acceptable salt, pharmaceutically acceptable N-oxide,
 pharmaceutically active metabolite, pharmaceutically acceptable prodrug, isomer
 derivative, or pharmaceutically acceptable solvate thereof.

Provided herein are compositions and methods for treating a disease comprising
 30 administering to a subject in need thereof an effective amount of a kinase modulating
 compound having the structure:

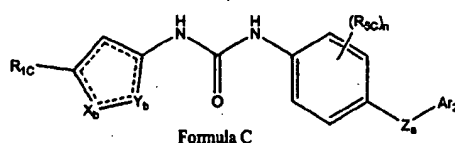


wherein:

- (a) X_b and Y_b are independently selected from O, N, NR_{c1} , and CR_c , wherein R_{c1} is a suitable substituent selected from hydrogen; alkyl, alkenyl, alkynyl, aryl, cycloalkyl, heterocycloalkyl, or heteroaryl unsubstituted or substituted with one, two, or three suitable substituents, wherein X_b and Y_b are not both oxygen;
- (b) R_{1b} and R_{2b} are each a suitable substituent independently selected from hydrogen, halogen, $=O$; $=S$; $-CN$; $-NO_2$, or an alkyl, alkenyl, heteroalkyl, haloalkyl, alkynyl, aryl, cycloalkyl, heterocycloalkyl, heteroaryl group unsubstituted or substituted with one, two or three suitable substituents independently selected from the group consisting of: halogens; $=O$; $=S$; $-CN$; and $-NO_2$; and alkyl, alkenyl, heteroalkyl, haloalkyl, alkynyl, aryl, cycloalkyl, heterocycloalkyl, heteroaryl, $-(CH_2)_zCN$ where z is a whole integer from 0 to 4, $=NH$, $-NHOH$, $-OH$, $-C(O)H$, $-OC(O)H$, $-C(O)OH$, $-OC(O)OH$, $-OC(O)OC(O)H$, $-OOH$, $-C(NH)NH_2$, $-NHC(NH)NH_2$, $-C(S)NH_2$, $-NHC(S)NH_2$, $-NHC(O)NH_2$, $-S(O_2)H$, $-S(O)H$, $-NH_2$, $-C(O)NH_2$, $-OC(O)NH_2$, $-NHC(O)H$, $-NHC(O)OH$, $-C(O)NHC(O)H$, $-OS(O_2)H$, $-OS(O)H$, $-OSH$, $-SC(O)H$, $-S(O)C(O)OH$, $-SO_2C(O)OH$, $-NHSH$, $-NHS(O)H$, $-NHHSO_2H$, $-C(O)SH$, $-C(O)S(O)H$, $-C(O)S(O_2)H$, $-C(S)H$, $-C(S)OH$, $-C(SO)OH$, $-C(SO_2)OH$, $-NHC(S)H$, $-OC(S)H$, $-OC(S)OH$, $-OC(SO_2)H$, $-S(O_2)NH_2$, $-S(O)NH_2$, $-SNH_2$, $-NHCS(O_2)H$, $-NHC(SO)H$, $-NHC(S)H$, and $-SH$ groups unsubstituted or substituted with one, two or three suitable substituents independently selected from the group consisting of halogens, $=O$, $-NO_2$, $-CN$, $-(CH_2)_zCN$ where z is a whole integer from 0 to 4, $-OR_c$, $-NR_cOR_c$, $-NR_cR_c$, $-C(O)NR_c$, $-C(O)OR_c$, $-C(O)R_c$, $-NR_cC(O)NR_cR_c$, $-NR_cC(O)R_c$, $-OC(O)OR_c$, $-OC(O)NR_cR_c$, $-SR_c$, unsubstituted alkyl, unsubstituted alkenyl, unsubstituted alkynyl, unsubstituted aryl, unsubstituted cycloalkyl, unsubstituted heterocycloalkyl, and unsubstituted heteroaryl, or two or more substituents cyclize to form a fused or spiro polycyclic cycloalkyl, heterocycloalkyl, aryl, or heteroaryl group, where each R_c is independently selected from hydrogen, unsubstituted alkyl, unsubstituted alkenyl, unsubstituted alkynyl, unsubstituted aryl, unsubstituted

cycloalkyl, unsubstituted heterocycloalkyl, and unsubstituted heteroaryl, or two or more R_c groups together cyclize to form part of a heteroaryl or heterocycloalkyl group unsubstituted or substituted with an unsubstituted alkyl group, or a pharmaceutically acceptable salt, pharmaceutically acceptable N-oxide, pharmaceutically active metabolite, pharmaceutically acceptable prodrug, or pharmaceutically acceptable solvate thereof.

Provided herein are compositions and methods for treating a disease by administering an effective amount of kinase modulating compound having the structure:

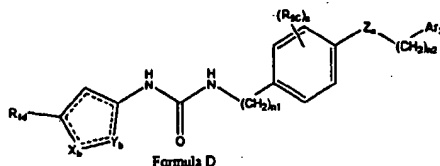


wherein:

- (a) R_{1C} is unsubstituted C_1 - C_5 alkyl or unsubstituted C_3 - C_6 cycloalkyl;
- (b) n is 0, 1 or 2; and
- (c) Each R_{5C} is a suitable substituent independently selected from the group consisting of halogens; -CN; and -NO₂; and alkyl, alkenyl, heteroalkyl, haloalkyl, alkynyl, aryl, cycloalkyl, heterocycloalkyl, heteroaryl, $-(CH_2)_zCN$ where z is a whole integer from 0 to 4, NH, -NHOH, -OH, -C(O)H, -OC(O)H, -C(O)OH, -OC(O)OH, -OC(O)OC(O)H, -OOH, -C(NH)NH₂, -NHC(NH)NH₂, -C(S)NH₂, -NHC(S)NH₂, -NHC(O)NH₂, -S(O₂)H, -S(O)H, -NH₂, -C(O)NH₂, -OC(O)NH₂, -NHC(O)H, -NHC(O)OH, -C(O)NHC(O)H, -OS(O₂)H, -OS(O)H, -OSH, -SC(O)H, -S(O)C(O)OH, -SO₂C(O)OH, -NHSH, -NHS(O)H, -NH₂SO₂H, -C(O)SH, -C(O)S(O)H, -C(O)S(O₂)H, -C(S)H, -C(S)OH, -C(SO)OH, -C(SO₂)OH, -NHC(S)H, -OC(S)H, -OC(S)OH, -OC(SO₂)H, -S(O₂)NH₂, -S(O)NH₂, -SNH₂, -NHCS(O₂)H, -NHC(SO)H, -NHC(S)H, and -SH groups unsubstituted or substituted with one, two or three suitable substituents independently selected from the group consisting of halogens, =O, -NO₂, -CN, $-(CH_2)_zCN$ where z 0, 1, 2, 3, or 4, -OR_c, -NR_cOR_c, -NR_cR_c, -C(O)NR_c, -C(O)OR_c, -C(O)R_c, -NR_cC(O)NR_cR_c, -NR_cC(O)R_c, -OC(O)OR_c, -OC(O)NR_cR_c, -SR_c, unsubstituted alkyl, unsubstituted alkenyl, unsubstituted alkynyl, unsubstituted aryl, unsubstituted cycloalkyl, unsubstituted heterocycloalkyl, and unsubstituted heteroaryl, or two or more substituents cyclize to form a fused or spiro polycyclic cycloalkyl, heterocycloalkyl, aryl,

or heteroaryl group, where each R_c is independently selected from hydrogen, unsubstituted alkyl, unsubstituted alkenyl, unsubstituted alkynyl, unsubstituted aryl, unsubstituted cycloalkyl, unsubstituted heterocycloalkyl, and unsubstituted heteroaryl, or two or more R_c groups together cyclize to form part of a heteroaryl or heterocycloalkyl group unsubstituted or substituted with an unsubstituted alkyl group.

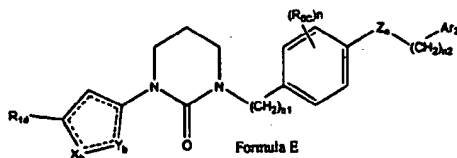
Provided herein are compositions and methods for treating a disease by administering an effective amount of kinase modulating compound having the structure:



wherein:

- (a) R_{1d} is unsubstituted C_1 - C_5 alkyl or unsubstituted C_3 - C_5 cycloalkyl;
- (b) n is 0, 1 or 2;
- (c) n_1 is 0, 1 or 2; and wherein n_2 is 0, 1 or 2; wherein n_1 and n_2 are not both 0.

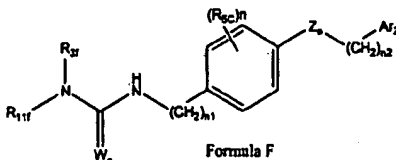
Provided herein are compositions and methods for treating a disease by administering an effective amount of a kinase modulating compound having the structure:



wherein:

- (a) n is 0, 1 or 2;

Provided herein are compositions and methods for treating a disease by administering an effective amount of a kinase modulating compound having the structure:

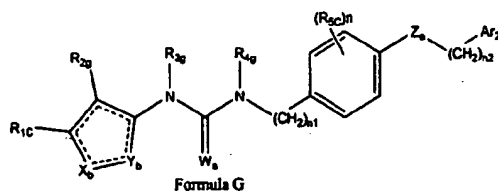


wherein:

- (a) R_{3f} and R_{11f} cyclize to form a heteroaryl or heterocycloalkyl group substituted or unsubstituted with one, two or three suitable substituents selected from the group

consisting of halogen, =O; =S; -CN; -NO₂, or an alkyl, alkenyl, heteroalkyl, haloalkyl, alkynyl, aryl, cycloalkyl, heterocycloalkyl, heteroaryl group unsubstituted or substituted with one, two or three suitable substituents independently selected from the group consisting of: halogens; =O; =S; -CN; and -NO₂; and alkyl, alkenyl, heteroalkyl, haloalkyl, alkynyl, aryl, cycloalkyl, heterocycloalkyl, heteroaryl, -(CH₂)_zCN where z is a whole integer from 0 to 4, =NH, -NHOH, -OH, -C(O)H, -OC(O)H, -C(O)OH, -OC(O)OH, -OC(O)OC(O)H, -OOH, -C(NH)NH₂, -NHC(NH)NH₂, -C(S)NH₂, -NHC(S)NH₂, -NHC(O)NH₂, -S(O₂)H, -S(O)H, -NH₂, -C(O)NH₂, -OC(O)NH₂, -NHC(O)H, -NHC(O)OH, -C(O)NHC(O)H, -OS(O₂)H, -OS(O)H, -OSH, -SC(O)H, -S(O)C(O)OH, -SO₂C(O)OH, -NHSH, -NHS(O)H, -NHSO₂H, -C(O)SH, -C(O)S(O)H, -C(O)S(O₂)H, -C(S)H, -C(S)OH, -C(SO)OH, -C(SO₂)OH, -NHC(S)H, -OC(S)H, -OC(S)OH, -OC(SO₂)H, -S(O₂)NH₂, -S(O)NH₂, -SNH₂, -NHCS(O₂)H, -NHC(SO)H, -NHC(S)H, and -SH groups unsubstituted or substituted with one, two or three suitable substituents independently selected from the group consisting of halogens, =O, -NO₂, -CN, -(CH₂)_zCN where z is a whole integer from 0 to 4, -OR_c, -NR_cOR_c, -NR_cR_c, -C(O)NR_c, -C(O)OR_c, -C(O)R_c, -NR_cC(O)NR_cR_c, -NR_cC(O)R_c, -OC(O)OR_c, -OC(O)NR_cR_c, -SR_c, unsubstituted alkyl, unsubstituted alkenyl, unsubstituted alkynyl, unsubstituted aryl, unsubstituted cycloalkyl, unsubstituted heterocycloalkyl, and unsubstituted heteroaryl, or two or more substituents cyclize to form a fused or spiro polycyclic cycloalkyl, heterocycloalkyl, aryl, or heteroaryl group, where each R_c is independently selected from hydrogen, unsubstituted alkyl, unsubstituted alkenyl, unsubstituted alkynyl, unsubstituted aryl, unsubstituted cycloalkyl, unsubstituted heterocycloalkyl, and unsubstituted heteroaryl, or two or more R_c groups together cyclize to form part of a heteroaryl or heterocycloalkyl group unsubstituted or substituted with an unsubstituted alkyl group.

Provided herein are compositions and methods for treating a disease comprising administering to a subject in need thereof an effective amount of a kinase modulating compound having the following structure:



wherein:

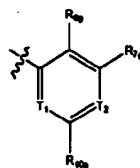
(a) R_{2g} , R_{3g} and R_{4g} are each independently selected from hydrogen, unsubstituted alkyl, unsubstituted aryl, and unsubstituted heteroaryl;

(b) n is 0, 1 or 2;

(c) n_1 is 0, 1 or 2;

(d) n_2 is 0, 1 or 2;

(e) Ar_2 is:



wherein:

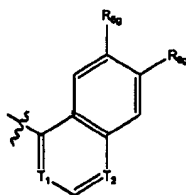
(i) R_{6g} and R_{7g} cyclize to form a 5- or 6-membered aryl, heteroaryl, heterocycloalkyl or cycloalkyl group unsubstituted or substituted with one, two or three suitable substituents independently selected from the group consisting of: halogens; -CN; and -NO₂; and alkyl, alkenyl, heteroalkyl, haloalkyl, alkynyl, aryl, cycloalkyl, heterocycloalkyl, heteroaryl, -(CH₂)_zCN where z is a whole integer from 0 to 4, NH, -NHOH, -OH, -C(O)H, -OC(O)H, -C(O)OH, -OC(O)OH, -OC(O)OC(O)H, -OOH, -C(NH)NH₂, -NHC(NH)NH₂, -C(S)NH₂, -NHC(S)NH₂, -NHC(O)NH₂, -S(O₂)H, -S(O)H, -NH₂, -C(O)NH₂, -OC(O)NH₂, -NHC(O)H, -NHC(O)OH, -C(O)NHC(O)H, -OS(O₂)H, -OS(O)H, -OSH, -SC(O)H, -S(O)C(O)OH, -SO₂C(O)OH, -NHSH, NHS(O)H, -NHSO₂H, -C(O)SH, -C(O)S(O)H, -C(O)S(O₂)H, -C(S)H, -C(S)OH, -C(SO)OH, -C(SO₂)OH, -NHC(S)H, -OC(S)H, -OC(S)OH, -OC(SO₂)H, -S(O₂)NH₂, -S(O)NH₂, -SNH₂, -NHCS(O₂)H, -NHC(SO)H, -NHC(S)H, and -SH groups unsubstituted or substituted with one, two or three suitable substituents independently selected from the group consisting of halogens, =O, -NO₂, -CN, -(CH₂)_z-CN where z is a whole integer from 0 to 4, -OR_c, -NR_cOR_c, -NR_cR_c, -C(O)NR_c, -C(O)OR_c, -C(O)R_c, -NR_cC(O)NR_cR_c,

- NR_cC(O)R_c, -OC(O)OR_c, -OC(O)NR_cR_c, -SR_c, unsubstituted alkyl, unsubstituted alkenyl, unsubstituted alkynyl, unsubstituted aryl, unsubstituted cycloalkyl, unsubstituted heterocycloalkyl, and unsubstituted heteroaryl, or two or more substituents cyclize to form a fused or spiro polycyclic cycloalkyl, heterocycloalkyl, aryl, or heteroaryl group, where each R_c is independently selected from hydrogen, unsubstituted alkyl, unsubstituted alkenyl, unsubstituted alkynyl, unsubstituted aryl, unsubstituted cycloalkyl, unsubstituted heterocycloalkyl, and unsubstituted heteroaryl, or two or more R_c groups together cyclize to form part of a heteroaryl or heterocycloalkyl group unsubstituted or substituted with an unsubstituted alkyl group;
- (ii) R_{10g} is a suitable substituent selected from hydrogen; halogens; -CN; and -NO₂; and alkyl, alkenyl, heteroalkyl, haloalkyl, alkynyl, aryl, cycloalkyl, heterocycloalkyl, heteroaryl, -(CH₂)_zCN where z is a whole integer from 0 to 4, NH, -NHOH, -OH, -C(O)H, -OC(O)H, -C(O)OH, -OC(O)OH, -OC(O)OC(O)H, -OOH, -C(NH)NH₂, -NHC(NH)NH₂, -C(S)NH₂, -NHC(S)NH₂, -NHC(O)NH₂, -S(O₂)H, -S(O)H, -NH₂, -C(O)NH₂, -OC(O)NH₂, -NHC(O)H, -NHC(O)OH, -C(O)NHC(O)H, -OS(O₂)H, -OS(O)H, -OSH, -SC(O)H, -S(O)C(O)OH, -SO₂C(O)OH, -NHSH, NHS(O)H, -NHSO₂H, -C(O)SH, -C(O)S(O)H, -C(O)S(O₂)H, -C(S)H, -C(S)OH, -C(SO)OH, -C(SO₂)OH, -NHC(S)H, -OC(S)H, -OC(S)OH, -OC(SO₂)H, -S(O₂)NH₂, -S(O)NH₂, -SNH₂, -NHCS(O₂)H, -NHC(SO)H, -NHC(S)H, and -SH groups unsubstituted or substituted with one, two or three suitable substituents independently selected from the group consisting of halogens, =O, -NO₂, -CN, -(CH₂)_zCN where z is a whole integer from 0 to 4, -OR_c, -NR_cOR_c, -NR_cR_c, -C(O)NR_c, -C(O)OR_c, -C(O)R_c, -NR_cC(O)NR_cR_c, -NR_cC(O)R_c, -OC(O)OR_c, -OC(O)NR_cR_c, -SR_c, unsubstituted alkyl, unsubstituted alkenyl, unsubstituted alkynyl, unsubstituted aryl, unsubstituted cycloalkyl, unsubstituted heterocycloalkyl, and unsubstituted heteroaryl, or two or more substituents cyclize to form a fused or spiro polycyclic cycloalkyl, heterocycloalkyl, aryl, or heteroaryl group, where each R_c is independently selected from hydrogen, unsubstituted alkyl, unsubstituted alkenyl, unsubstituted alkynyl, unsubstituted aryl, unsubstituted cycloalkyl, unsubstituted heterocycloalkyl, and unsubstituted heteroaryl, or two or more R_c groups together cyclize to form part of a heteroaryl or heterocycloalkyl group unsubstituted or substituted with an unsubstituted alkyl group; and

(iii) T_1 and T_2 are each independently selected from CR_w and N, where R_w is a suitable substituent selected from hydrogen; halogens; -CN; and -NO₂; and unsubstituted alkyl, unsubstituted alkenyl, unsubstituted alkynyl, unsubstituted aryl, unsubstituted cycloalkyl, unsubstituted heterocycloalkyl, and unsubstituted heteroaryl, or two or more substituents cyclize to form a fused or spiro polycyclic cycloalkyl, heterocycloalkyl, aryl, or heteroaryl group;

or a pharmaceutically acceptable salt, pharmaceutically acceptable N-oxide, pharmaceutically active metabolite, pharmaceutically acceptable prodrug, or pharmaceutically acceptable solvate thereof.

Compositions and methods of Formulas A-G are provided wherein X_b is O and Y_b is N and/or X_b is N and Y_b is O; and/or R_{2a} , R_{2g} , R_{3a} , R_{3g} , R_{4a} and R_{4g} are each hydrogen; and/or R_{1b} , R_{1c} , and R_{1d} are each an unsubstituted or substituted t-butyl and R_{2b} and R_{2g} are hydrogen; and/or W_a is O; and/or Z_a is C(O)NH or NHC(O); and/or n is 0. In various embodiments, T_1 is N and T_2 is N or T_1 is N and T_2 is CH. In other embodiments, Ar_2 is:



wherein:

(i) R_{8g} and R_{9g} are suitable substituents each independently selected from the group consisting of hydrogen; halogens; -CN; and -NO₂; and alkyl, alkenyl, heteroalkyl, haloalkyl, alkynyl, aryl, cycloalkyl, heterocycloalkyl, heteroaryl, -(CH₂)_zCN where z is a whole integer from 0 to 4, -NH, -NHOH, -OH, -C(O)H, -OC(O)H, -C(O)OH, -OC(O)OH, -OC(O)OC(O)H, -OOH, -C(NH)NH₂, -NHC(NH)NH₂, -C(S)NH₂, -NHC(S)NH₂, -NHC(O)NH₂, -S(O₂)H, -S(O)H, -NH₂, -C(O)NH₂, -OC(O)NH₂, -NHC(O)H, -NHC(O)OH, -C(O)NHC(O)H, -OS(O₂)H, -OS(O)H, -OSH, -SC(O)H, -S(O)C(O)OH, -SO₂C(O)OH, -NHSH, -NHS(O)H, -NH₂SO₂H, -C(O)SH, -C(O)S(O)H, -C(O)S(O₂)H, -C(S)H, -C(S)OH, -C(SO)OH, -C(SO₂)OH, -NHC(S)H, -OC(S)H, -OC(S)OH, -OC(SO₂)H, -S(O₂)NH₂, -S(O)NH₂, -SNH₂, -NHCS(O₂)H, -NHC(SO)H, -NHC(S)H, and -SH groups unsubstituted or substituted with one, two or three suitable substituents independently selected from the group consisting of halogens,

=O, -NO₂, -CN, -(CH₂)_z-CN where z is a whole integer from 0 to
 4, -OR_c, -NR_cOR_c, -NR_cR_c, -C(O)NR_c, -C(O)OR_c, -C(O)R_c, -NR_cC(O)NR_cR_c,
 -NR_cC(O)R_c, -OC(O)OR_c, -OC(O)NR_cR_c, -SR_c, unsubstituted alkyl, unsubstituted
 5 alkenyl, unsubstituted alkynyl, unsubstituted aryl, unsubstituted cycloalkyl, unsubstituted
 heterocycloalkyl, and unsubstituted heteroaryl, or two or more substituents cyclize to
 form a fused or spiro polycyclic cycloalkyl, heterocycloalkyl, aryl, or heteroaryl group,
 where each R_c is independently selected from hydrogen, unsubstituted alkyl,
 unsubstituted alkenyl, unsubstituted alkynyl, unsubstituted aryl, unsubstituted cycloalkyl,
 unsubstituted heterocycloalkyl, and unsubstituted heteroaryl, or two or more R_c groups
 10 together cyclize to form part of a heteroaryl or heterocycloalkyl group unsubstituted or
 substituted with an unsubstituted alkyl group; and
 (ii) T₁ is N and T₂ is CH or N.

Compositions and methods of Formulas A-G are provided herein wherein R_{8g} and R_{9g} are
 each independently selected from the group consisting of hydrogen; halogens; -CN;
 15 and -NO₂; and alkyl, alkenyl, heteroalkyl, haloalkyl, alkynyl, aryl, cycloalkyl,
 heterocycloalkyl, heteroaryl, -(CH₂)_zCN where z is a whole integer from 0 to
 4, -OH, -C(O)H, -OC(O)H, -C(O)OH, -NH₂, -C(O)NH₂, -NHC(O), -OC(O)NH₂, -NHC(O)H,
 -NHC(O)OH groups unsubstituted or substituted with one, two or three suitable substituents
 independently selected from the group consisting of halogens, unsubstituted alkyl,
 20 unsubstituted alkenyl, unsubstituted alkynyl, unsubstituted aryl, unsubstituted cycloalkyl,
 unsubstituted heterocycloalkyl, and unsubstituted heteroaryl, or two or more substituents
 cyclize to form a fused or spiro polycyclic cycloalkyl, heterocycloalkyl, aryl, or heteroaryl
 group.

Compositions and methods of Formulas A-G are provided herein wherein each R_{5C} is a
 25 suitable substituent independently selected from the group consisting of halogens; -CN;
 and -NO₂; and alkyl, alkenyl, heteroalkyl, haloalkyl, alkynyl, aryl, cycloalkyl,
 heterocycloalkyl, heteroaryl, -(CH₂)_zCN where z is a whole integer from 0 to
 4, -OH, -C(O)H, -OC(O)H, -C(O)OH, -NH₂, -C(O)NH₂, -NHC(O), -OC(O)NH₂,
 -NHC(O)H, -NHC(O)OH groups unsubstituted or substituted with one, two or three suitable
 30 substituents independently selected from the group consisting of halogens, unsubstituted
 alkyl, unsubstituted alkenyl, unsubstituted alkynyl, unsubstituted aryl, unsubstituted

cycloalkyl, unsubstituted heterocycloalkyl, and unsubstituted heteroaryl, or two or more substituents cyclize to form a fused or spiro polycyclic cycloalkyl, heterocycloalkyl, aryl, or heteroaryl group.

5 Compositions and methods of Formula A are provided herein wherein Ar₃ is a 5-membered aryl, heteroaryl, heterocycloalkyl or cycloalkyl group unsubstituted or substituted with one, two or three suitable substituents. In some embodiments, Ar₃ is a 5- or 6-membered aryl or heteroaryl group unsubstituted or substituted with one, two or three suitable substituents.

10 Compositions and methods of Formula A-G are provided herein wherein n₃ is 0 or 1, and/or wherein n₁ is 0, 1 or 2, and/or n₂ is 0, 1 or 2. In some embodiments, R_{3a}/R_{3g} and R_{4a}/R_{4g} are each hydrogen. In other embodiments, R_{3a}/R_{3g} and R_{4a}/R_{4g} are not both substituted.

15 Compositions and methods of Formula A are provided herein wherein Ar₃ is a substituted or unsubstituted 5-membered heteroaryl group and R₂ is hydrogen. In some embodiments, Ar₁ is an unsubstituted or substituted 6-membered aryl group or an unsubstituted or substituted 6-membered heteroaryl group. In other embodiments, W_a is O.

20 Compositions and methods of Formulas A-G wherein Z_a is not carbon are described herein. In some embodiments, Z_a is selected from S, O, N, NR_{c2}, C(O)NR_{c2}, and NR_{c2}C(O), wherein R_{c2} is hydrogen, unsubstituted alkyl, unsubstituted alkenyl, unsubstituted alkynyl, unsubstituted aryl, unsubstituted cycloalkyl, unsubstituted heterocycloalkyl, or unsubstituted heteroaryl group. In other embodiments, Z_a is C(O)NH, NHC(O), or NH.

Compositions and methods of Formulas A-G wherein W_a is S, O, or NH are described herein.

25 Compositions and methods of Formulas A-G are described herein wherein Ar₁ is an aryl or heteroaryl group unsubstituted or substituted with one, two or three suitable substituents independently selected from the group consisting of halogens; -CN; and -NO₂; and alkyl, alkenyl, heteroalkyl, haloalkyl, alkynyl, aryl, cycloalkyl, heterocycloalkyl, heteroaryl, -(CH₂)_zCN where z is a whole integer from 0 to 4, -OH, -C(O)H, -OC(O)H, -C(O)OH, -NH₂, -C(O)NH₂, -NHC(O), -OC(O)NH₂, -NHC(O)H, -NHC(O)OH groups
30 unsubstituted or substituted with one, two or three suitable substituents independently selected from the group consisting of halogens, unsubstituted alkyl, unsubstituted alkenyl,

unsubstituted alkynyl, unsubstituted aryl, unsubstituted cycloalkyl, unsubstituted heterocycloalkyl, and unsubstituted heteroaryl.

Compositions and methods of Formulas A-G are described herein wherein Ar₂ is an aryl, heteroaryl, heterocycloalkyl or cycloalkyl group unsubstituted or substituted with one, two or three suitable substituents independently selected from the group consisting of
5 halogens; -CN; and -NO₂; and alkyl, alkenyl, heteroalkyl, haloalkyl, alkynyl, aryl, cycloalkyl, heterocycloalkyl, heteroaryl, -(CH₂)_zCN where z is a whole integer from 0 to 4, -OH, -C(O)H, -OC(O)H, -C(O)OH, -NH₂, -C(O)NH₂, -NHC(O), -OC(O)NH₂, -NHC(O)H, -NHC(O)OH groups unsubstituted or substituted with one, two or three suitable substituents
10 independently selected from the group consisting of halogens, unsubstituted alkyl, unsubstituted alkenyl, unsubstituted alkynyl, unsubstituted aryl, unsubstituted cycloalkyl, unsubstituted heterocycloalkyl, and unsubstituted heteroaryl. In some embodiments, Ar₂ is an unsubstituted or substituted pyridinyl. In other embodiments, Ar₂ is an unsubstituted or substituted quinazolinyl.

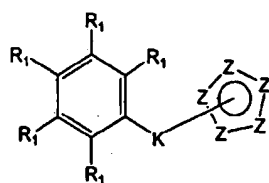
Compositions and methods of Formulas A-G are provided herein wherein X_b and Y_b are each independently selected from O, N, and NR_{c1} wherein R_{c1} is unsubstituted alkyl or unsubstituted aryl. In some embodiments, X_b is N and Y_b is NR_{c1}. In other embodiments, X_b is O and Y_b is N. In other embodiments, X_b is O and Y_b is N, or X_b is N and Y_b is O.

Compositions and methods of Formulas A-G are provided herein wherein R₁ is
20 unsubstituted t-butyl or unsubstituted cyclopropyl.

Compositions and methods are provided herein wherein R_{5c} is independently selected from the group consisting of halogens, -CN, -NO₂, unsubstituted alkyl, unsubstituted alkenyl, unsubstituted heteroalkyl, unsubstituted haloalkyl, unsubstituted alkynyl, unsubstituted aryl, unsubstituted cycloalkyl, unsubstituted heterocycloalkyl, and unsubstituted heteroaryl group.

Compositions and methods of Formulas A-G are provided herein wherein R_{10g} is
25 hydrogen or lower alkyl.

Provided herein are compositions and methods for treating a disease comprising administering to a subject in need thereof an effective amount of a kinase modulating compound having the following structure:



IA

wherein:

each Z is independently C, CR₃, N, NR₃, O, or S, provided that no more than two

Z's are heteroatoms and wherein no two adjacent Z's are O or S,

where R₃ is H, substituted or unsubstituted alkyl, substituted or unsubstituted cycloalkyl, substituted or unsubstituted heteroaryl, or substituted or unsubstituted aryl; and

each R₁ is independently H, halogen, substituted or unsubstituted alkyl,

substituted or unsubstituted alkoxy, substituted or unsubstituted cycloalkyl,

substituted or unsubstituted heterocyclyl, substituted or unsubstituted aryl,

substituted or unsubstituted heteroaryl, -OR_c, -OC(O)R_c, -NO₂, -N(R_c)₂, -SR_c,

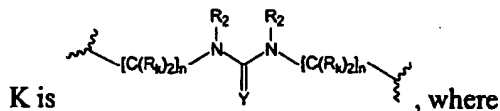
S(O)_jR_c where j is 1 or 2, -NR_cC(O)R_c, -C(O)N(R_c)₂, -C(O)₂R_c, or -C(O)R_c; or

two adjacent R₁'s, are taken together to form a substituted or unsubstituted aryl or heteroaryl, where

each R_c is independently H, substituted or unsubstituted alkyl, substituted

or unsubstituted cycloalkyl, substituted or unsubstituted aryl, or

substituted or unsubstituted heteroaryl.



Y is O or S;

each R_k is independently H, halogen, substituted or unsubstituted

alkyl, -OR₂, substituted or unsubstituted alkoxy, -OC(O)R₂, -

NO₂, -N(R₂)₂,

-SR₂, -C(O)R₂, -C(O)₂R₂, -C(O)N(R₂)₂, or -N(R₂)C(O)R₂;

each R₂ is independently H, substituted or unsubstituted alkyl, substituted

or unsubstituted cycloalkyl, substituted or unsubstituted heterocyclyl,

substituted or unsubstituted aryl, or substituted or unsubstituted

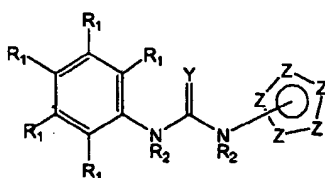
heteroaryl; or wherein two R_2 groups are linked together by an optionally substituted alkylene; and

each n is independently 0, 1, 2, 3 or 4;

or an active metabolite, or a pharmaceutically acceptable prodrug, isomer,

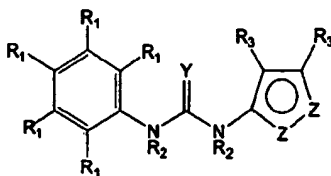
5 pharmaceutically acceptable salt or solvate thereof.

Provided herein are compositions and methods for treating a disease comprising administering to a subject in need thereof an effective amount of a kinase modulating compound having the following structure:



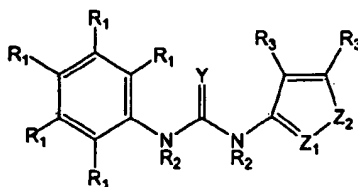
10 (I).

Provided herein are compositions and methods for treating a disease comprising administering to a subject in need thereof an effective amount of a kinase modulating compound having the following structure:



15 (II).

Provided herein are compositions and methods for treating a disease comprising administering to a subject in need thereof an effective amount of a kinase modulating compound having the following structure:



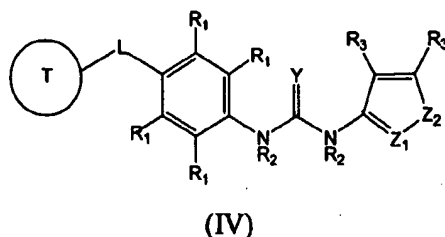
20 (III)

wherein:

Z_1 is CR_3 or N; and

Z_2 is O or S.

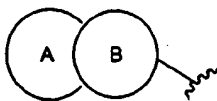
Provided herein are compositions and methods for treating a disease comprising administering to a subject in need thereof an effective amount of a kinase modulating compound having the following structure:



wherein:

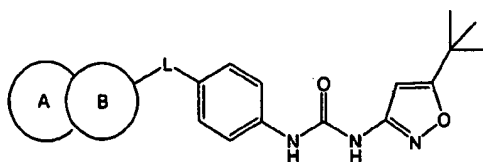
L is a linker selected from the group consisting of a covalent bond, -(substituted or unsubstituted alkylene)-, -(substituted or unsubstituted alkenylene)-, -O-, -O(substituted or unsubstituted alkylene)-, -C(O)-, -C(O)(substituted or unsubstituted alkylene)-, -C(O)(substituted or unsubstituted alkenylene)-, -NH-, -NH(substituted or unsubstituted alkylene)-, -NH(substituted or unsubstituted alkenylene)-, -C(O)NH-, -C(O)NH(substituted or unsubstituted alkylene)-, -C(O)NH(substituted or unsubstituted alkenylene)-, -NHC(O)(substituted or unsubstituted alkylene)-, -NHC(O)(substituted or unsubstituted alkenylene)-, -S-, -S(substituted or unsubstituted alkylene)-, and -NHC(O)(substituted or unsubstituted alkylene)S(substituted or unsubstituted alkylene)C(O)NH-; and

T is a mono-, bi-, or tricyclic, substituted or unsubstituted cycloalkyl, heterocyclyl, aryl, or heteroaryl.



In some embodiments, T is wherein A is a substituted or unsubstituted five or six-membered aryl, heterocyclyl or heteroaryl; and B is a substituted or unsubstituted five or six-membered arylene, heterocyclene or heteroarylene, wherein A and B together form a fused two-ring moiety.

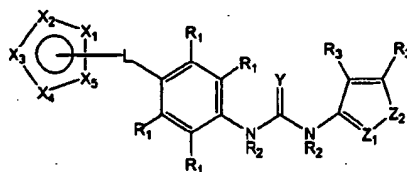
Provided herein are compositions and methods for treating a disease comprising administering to a subject in need thereof an effective amount of a kinase modulating compound having the following structure:



(VI).

In some embodiments, L is $-\text{C}(\text{O})\text{NH}-$. In other embodiments, B is substituted or unsubstituted phenylene, pyridinylene, pyrimidinylene, pyridazinylene, thiophenylene, imidazolylene, or pyrrolylene. In still other embodiments, L is $-\text{NH}-$. In yet other embodiments, B is substituted or unsubstituted phenylene, pyridinylene, pyrimidinylene, pyridazinylene, thiophenylene, or imidazolylene.

Provided herein are compositions and methods for treating a disease comprising administering to a subject in need thereof an effective amount of a kinase modulating compound having the following structure:



(VII)

wherein:

each of $\text{X}_1\text{-X}_5$ is independently C, CR, N, NR, S, or O, wherein no more than three of

$\text{X}_1\text{-X}_5$ is a heteroatom, and no two adjacent ring atoms are O or S; where

each R is independently H, halogen, substituted or unsubstituted alkyl, $-\text{OH}$,

substituted or unsubstituted alkoxy, $-\text{OC}(\text{O})\text{R}_d$, $-\text{NO}_2$, $-\text{N}(\text{R}_d)_2$, $-\text{SR}_d$, $-\text{S}(\text{O})_j\text{R}_d$

where j is 1 or 2, $-\text{NR}_d\text{C}(\text{O})\text{R}_d$, $-\text{C}(\text{O})_2\text{R}_d$, $-\text{C}(\text{O})\text{N}(\text{R}_d)_2$, or $-\text{C}(\text{O})\text{R}_d$, or two

adjacent R_d 's are taken together to form a substituted or unsubstituted aryl or

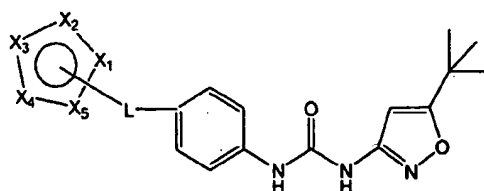
heteroaryl,

where each R_d is independently H, substituted or unsubstituted alkyl,

substituted or unsubstituted cycloalkyl, substituted or unsubstituted

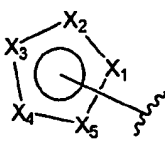
aryl, or substituted or unsubstituted heteroaryl.

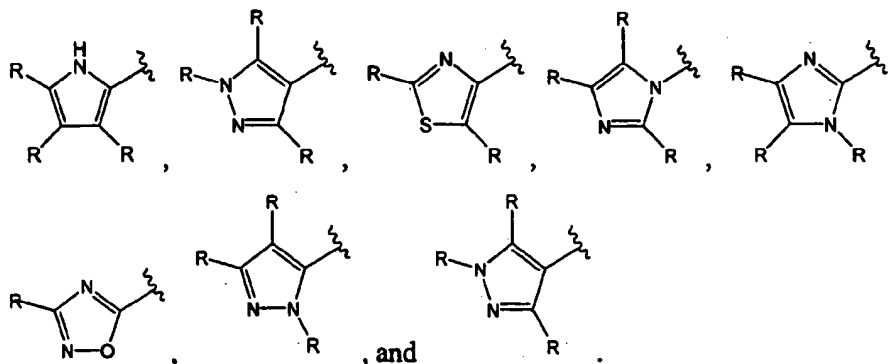
Provided herein are compositions and methods for treating a disease comprising administering to a subject in need thereof an effective amount of a kinase modulating compound having the following structure:



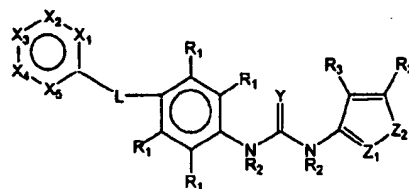
(VIII).

In some embodiments, L is a covalent bond, $-C(O)NH-$, $-OCH_2-$, or $-OCH_2CH_2-$. In other

embodiments,  is selected from the group consisting of:

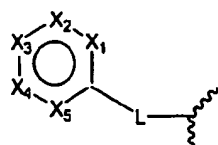


Provided herein are compositions and methods for treating a disease comprising administering to a subject in need thereof an effective amount of a kinase modulating compound having the following structure:



(IX)

wherein:



is selected from the group consisting of:

- (a) L is selected from the group consisting of $-O$ (substituted or unsubstituted alkylene)-, and $-C(O)$ (substituted or unsubstituted alkenylene)-; and

each of X_1 - X_5 is independently CR, N-O, or N, wherein no more than two of X_1 - X_5 is N, where

each R is independently H, halogen, substituted or unsubstituted alkyl, -OH, substituted or unsubstituted alkoxy, -OC(O) R_d , -NO₂, -N(R_d)₂, -SR_d, -NR_dC(O) R_d , or -C(O) R_d ,

each R_d is independently H, substituted or unsubstituted alkyl, substituted or unsubstituted cycloalkyl, substituted or unsubstituted aryl, or substituted or unsubstituted heteroaryl;

(b) L is -NH-;

each of X_1 , X_2 , X_4 , and X_5 is independently CR, N-O, or N; and X_3 is independently CR₅ or N, wherein no more than two of X_1 - X_5 is N, where

R_5 is selected from the group consisting of H, halogen, substituted or unsubstituted alkyl, substituted alkoxy, -C(O) R_d , -OC(O) R_d , -NO₂, -N(R_d)₂, and -SR_d, and

each R_d is independently H, substituted or unsubstituted alkyl, substituted or unsubstituted cycloalkyl, substituted or unsubstituted aryl, or substituted or unsubstituted heteroaryl;

(c) L is -NH-;

each of X_1 , X_3 , and X_5 is independently CR, N-O, or N; and

each of X_2 and X_4 is independently CR₆ or N, wherein no more than two of X_1 - X_5 is N; where

R_6 is selected from the group consisting of H, halogen, unsubstituted alkyl, -OH, substituted or unsubstituted alkoxy, -C(O) R_d , -OC(O) R_d ,

-NO₂, -N(R_d)₂, -SR_d, and alkyl substituted with alkoxy, halogen, aryl, heteroaryl, amine, -C(O) R_d , -OC(O) R_d , -NO₂, -N(R_d)₂, and -SR_d, and

each R_d is independently H, substituted or unsubstituted alkyl, substituted or unsubstituted cycloalkyl, substituted or unsubstituted aryl, or substituted or unsubstituted heteroaryl;

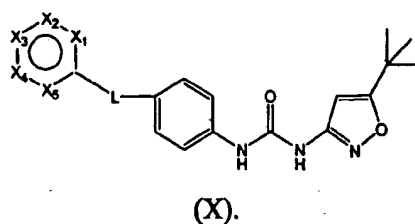
(d) L is -C(O)NH-;

each of X_1 , X_2 , X_4 , and X_5 is independently CR, N-O, or N; and
 X_3 is independently CR₇ or N, wherein no more than two of X_1 - X_5 is N, and
 when X_3 is N, at least one of X_1 , X_2 , X_3 , or X_5 is not CH, where

R_7 is selected from the group consisting of H, halogen, substituted or
 unsubstituted alkyl, -OH, substituted or unsubstituted alkoxy,
 -C(O) R_d , -OC(O) R_d , -NO₂, -N(R_d)₂, and -SR_d, and

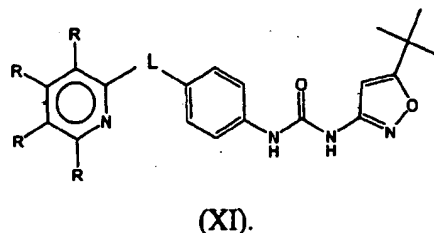
each R_d is independently H, substituted or unsubstituted alkyl,
 substituted or unsubstituted cycloalkyl, substituted or unsubstituted
 aryl, or substituted or unsubstituted heteroaryl.

Provided herein are compositions and methods for treating a disease comprising
 administering to a subject in need thereof an effective amount of a kinase modulating
 compound having the following structure:



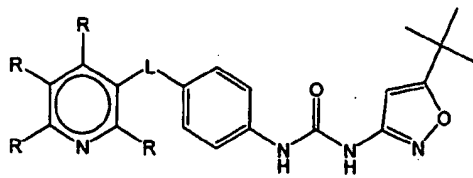
In some embodiments, L is a linker selected from the group consisting of a covalent bond, -
 (substituted or unsubstituted alkylene), -NHC(O)-, -C(O)NH(substituted or unsubstituted
 alkylene), -NHC(O)(substituted or unsubstituted alkylene), -C(O)NH(substituted or
 unsubstituted alkenylene), -NHC(O)(substituted or unsubstituted alkenylene)-, and -
 O(substituted or unsubstituted alkylene)-.

Provided herein are compositions and methods for treating a disease comprising
 administering to a subject in need thereof an effective amount of a kinase modulating
 compound having the following structure:



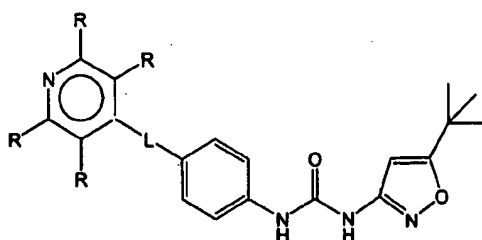
Provided herein are compositions and methods for treating a disease comprising
 administering to a subject in need thereof an effective amount of a kinase modulating

compound having the following structure:



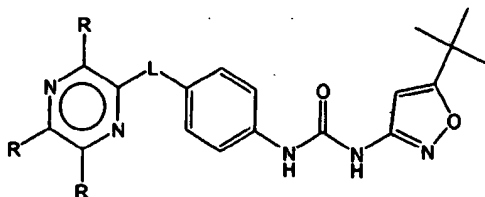
(XII).

5 Provided herein are compositions and methods for treating a disease comprising administering to a subject in need thereof an effective amount of a kinase modulating compound having the following structure:



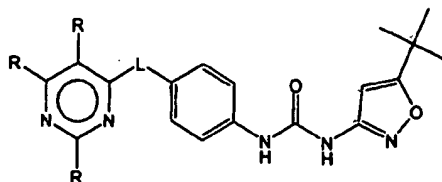
(XIII).

10 Provided herein are compositions and methods for treating a disease comprising administering to a subject in need thereof an effective amount of a kinase modulating compound having the following structure:



(XIV).

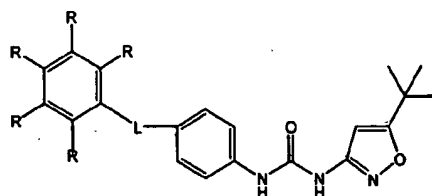
15 Provided herein are compositions and methods for treating a disease comprising administering to a subject in need thereof an effective amount of a kinase modulating compound having the following structure:



(XV).

Provided herein are compositions and methods for treating a disease comprising

administering to a subject in need thereof an effective amount of a kinase modulating compound having the following structure:

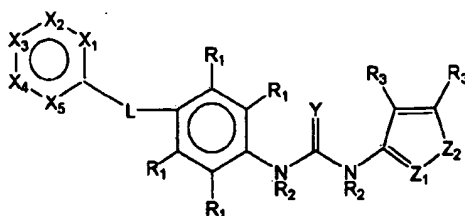


(XVI).

- 5 In some embodiments, L is a linker selected from the group consisting of -NHC(O)-, -OCH₂-, -OCH₂CH₂-, -NHC(O)CH₂SCH₂C(O)NH-, -CHCHC(O)NH-, -CHCHCH₂O-, -CH₂CH₂-, and -CH₂CH₂C(O)NH-.

Provided herein are compositions and methods for treating a disease comprising administering to a subject in need thereof an effective amount of a kinase modulating compound having the following structure:

10



(XVII)

wherein:

15

each of X₁-X₅ is independently C, CR, N-O, or, wherein no more than two of X₁-X₅ is N; where

each R is independently H, halogen, substituted or unsubstituted alkyl, -OR_d, substituted or unsubstituted alkoxy, -OC(O)R_d, -NO₂, -N(R_d)₂, -SR_d, -S(O)_jR_d where j is 1 or 2, -NR_d C(O)R_d, -C(O)₂R_d, -C(O)N(R_d)₂ or -C(O)R_d; or two adjacent R's are taken together to form a substituted or unsubstituted aryl or hetroaryl, and

20

each R_d is independently H, substituted or unsubstituted alkyl, substituted or unsubstituted cycloalkyl, substituted or unsubstituted aryl, or substituted or unsubstituted heteroaryl

with a proviso that said compound is not: